CLAIMS

What is claimed is:

1. A compound of the formula (I):

wherein:

W is:

wherein:

m is 0 or 1;

each R^1 is hydroxy, alkoxy, or aryloxy, or each R^1 is an oxygen atom and together with the boron, to which they are each bound, form a 5-7 membered ring,

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wherein the ring atoms are carbon, nitrogen, or oxygen;

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each R² is independently hydrogen, alkyl, alkenyl, aryl, aralkyl, aralkenyl, cycloalkyl, cycloalkylalkyl, cycloalkenyl, cycloalkenylalkyl, heterocyclyl, heterocyclylalkyl, heterocyclylalkenyl, heteroaryl, or heteroaralkyl, or two R² groups, which are bound to the same nitrogen atom, form together with that nitrogen atom, a 5-7 membered monocyclic heterocyclic ring system; wherein any R² carbon atom is optionally substituted with J;

J is alkyl, aryl, aralkyl, alkoxy, aryloxy, aralkoxy, cycloalkyl, cycloalkoxy, heterocyclyl, heterocyclyloxy, heterocyclylalkyl, keto, hydroxy, amino, alkylamino, alkanoylamino, aroylamino, aralkanoylamino, carboxy, carboxyalkyl, carboxamidoalkyl, halo, cyano, nitro, formyl, acyl, sulfonyl, or sulfonamido and is optionally substituted with 1-3 J¹ groups; and

J1 is alkyl, aryl, aralkyl, alkoxy, aryloxy, heterocyclyl, heterocyclyloxy, keto, hydroxy, amino, alkanoylamino, aroylamino, carboxy, carboxyalkyl, carboxamidoalkyl, halo, cyano, nitro, formyl, sulfonyl, or sulfonamido;

L is alkyl, alkenyl, or alkynyl, wherein any hydrogen is optionally substituted with halogen, and wherein any hydrogen or halogen atom bound to any terminal carbon atom is optionally substituted with sulfhydryl or hydroxy;

each M is independently alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, or

heteroaralkyl, and is optionally substituted by 1 to 3 J groups, wherein any alkyl carbon atom may be replaced by a heteroatom;

 R^{18} is a bond, $-N(R^{11})$ -, or -C(0)-; R^{11} is hydrogen or C_1 - C_3 alkyl;

each R^{19} is independently H or R^{21} -aryl, or 2 adjacent R^{19} may be bound to one another to form a 5-7 membered aromatic ring; wherein any R^{19} is optionally substituted with 1 to 4 independently selected J' groups;

each R^{21} is independently C_1-C_3 -straight or branched alkyl, C_2-C_3 -straight or branched alkenyl, $O-(C_1-C_3)$ -straight or branched alkyl, or $O-(C_2-C_3)$ -straight or branched alkenyl;

n is 0 or 1;

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the ring to which R^{18} and R^{19} are attached may be saturated, partially saturated, aromatic or fully unsaturated; and 1 to 3 carbon atoms that make up the ring to which R^{18} and R^{19} are attached are optionally replaced with a heteroatom which is independently selected from O, S, S(O), S(O)₂ or N(R^{11});

 A^2 is a bond or $-N(R^{11})-R^{17}(M)-R^{22}-$, wherein

 R^{17} is -CH- or -N-; and R^{22} is -C(0)- or -S(0)₂-;

V is a bond, $-CH(R^{11})$ -, -O -, -S -, or $-N(R^{11})$ -;

K is a bond, -0-, -S-, -C(0)-, -S(0)-, $-S(0)_2-$, or -

 $S(O) NR^{11}$ -; $T \text{ is } -R^{12}, \text{ } -\text{alkyl-}R^{12}, \text{ } -\text{alkenyl-}R^{12}, \text{ } -\text{alkynyl-}R^{12},$ $-OR^{12}, -N(R^{12})_2, -C(O)R^{12}, -C(=NO-\text{alkyl})R^{12}, \text{ or}$

$$R^{16}$$
 K N H O R^{10} , wherein:

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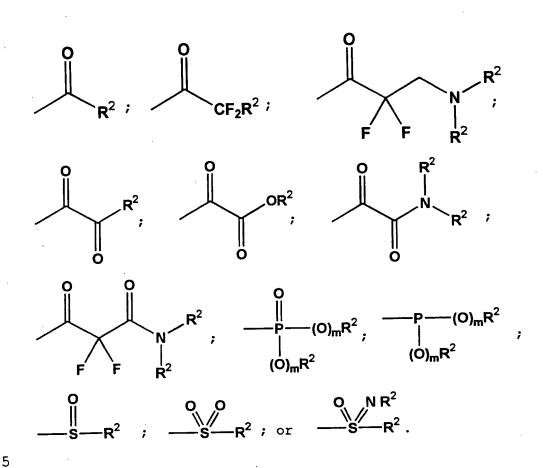
each R¹² is independently selected from hydrogen, aryl, heteroaryl, cycloalkyl, heterocyclyl, cycloalkylidenyl, or heterocycloalkylidenyl, and is optionally substituted with 1 to 3 J groups; or a first R¹² and a second R¹², together with the nitrogen to which they are bound, form a mono- or bicyclic ring system optionally substituted by 1 to 3 J groups;

R¹⁰ is alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, heteroaralkyl, carboxyalkyl, or carboxamidoalkyl, and is optionally substituted with 1 to 3 hydrogens J groups;

R¹⁵ is alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, heteroaralkyl, carboxyalkyl, or carboxamidoalkyl, and is optionally substituted with 1 to 3 J groups; and

 ${\ensuremath{\mathsf{R}}}^{16}$ is hydrogen, alkyl, aryl, heteroaryl, cycloalkyl, or heterocyclyl.

2. The compound according to claim 1, wherein W is selected from:



3. The compound according to claim 2, wherein W is -C(0)H.

4. The compound according to claim 1, wherein

10 J is selected from alkyl, alkoxy, aryloxy, aryl, aralkyl,
aralkoxy, halo, heteroaryl, cyano, amino, nitro,
heterocyclyl, acyl, carboxy, carboxyalkyl, alkylamino,
hydroxy, heterocyclylalkyl, aralkanoylamino, aroylamino,
alkanoylamino, formyl or keto.

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J is selected from t-butyl, methyl, trifluoromethyl, hydroxy, methoxy, ethoxy, trifluoromethoxy, carboxy, phenyl, benzyl, phenoxy, benzyloxy, fluoro, chloro, bromo, isoxazolyl, pyridinyl, piperidinyl, carboxymethyl, carboxyethyl, dialkylamino, morpholinylmethyl,

phenylacetylamino, or acylamino.

- 6. The compound according to claim 1, wherein each J^1 is independently selected from alkoxy, alkyl, halo or aryl.
 - 7. The compound according to claim 6, wherein each J^1 is independently selected from C_{1-3} alkoxy, chloro, C_{1-3} alkyl, or phenyl.
- 10 8. The compound according to claim 1, wherein L is selected from trihalomethyl, sulfhydryl or alkyl substituted with trihalomethyl, sulfhydryl, or hydroxy.
- 9. The compound according to claim 8, wherein 15 L is $-CH_2CH_3$ or $-CH_2CF_3$.
- 10. The compound according to claim 1, wherein R² is selected from H, fluorine, trifluoromethyl, alkyl, aryl, aralkyl, heteroaralkyl, heterocyclyl, or heterocyclylalkyl.
 - 11. The compound according to claim 10, wherein \mathbb{R}^2 is H.
- 25 12. The compound according to claim 1, wherein each M is independently selected from isopropyl, propyl, methyl, pyridylmethyl, benzyl, naphthylmethyl, phenyl, imidazolylmethyl, thiophenylmethyl, cyclohexylmethyl, phenethyl, benzylthiomethyl, or benzyloxyethyl.

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13. The compound according to claim 12, wherein each M is isopropyl.

- 14. The compound according to claim 1, wherein one R^{19} is R^{21} -aryl and the other two R^{19} are H, or two R^{19} are bound together to form an aromatic ring and the other R^{19} is H.
- 15. The compound according to claim 14, wherein one R^{19} is $-O-(C_1-C_3)$ -alkyl-aryl.

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- 16. The compound according to claim 15, wherein 14, wherein one \mathbb{R}^{19} is -O-benzyl.
- 17. The compound according to claim 14,

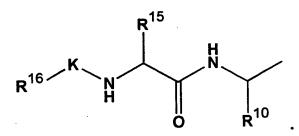
 15 wherein the two R¹⁹ that are bound together form a 6membered aromatic ring.
 - 18. The compound according to claim 17, wherein the two R^{19} that are bound together form phenyl.

- 19. The compound according to claim 1, wherein R^{18} is $-N\left(R^{11}\right)$ -.
- 20. The compound according to claim 19, wherein R^{18} is -N(H) or $-N(CH_3)$ -.
 - 21. The compound according to claim 1, wherein A^2 is a bond or $-N(R^{11})-C(M)-C(O)-$.
- 30 22. The compound according to claim 21, wherein A² is a bond or -N(H)-C(M)-C(O)-, wherein M is isopropyl.

- 23. The compound according to claim 1, wherein V is $-N(R^{11})$ -.
- 5 24. The compound according to claim 23, wherein V is -NH-.
 - 25. The compound according to claim 1, wherein K is -C(0) or $-S(0)_2$.

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- 26. The compound according to claim 25, wherein K is -C(0)-.
- 27. The compound according to claim 1, wherein 15 T is selected from $-R^{12}$, $-alkyl-R^{12}$, $-alkenyl-R^{12}$, $-OR^{12}$, $-N(R^{12})_2$, $-C(=NO-alkyl)-R^{12}$, or



- 28. The compound according to claim 27, wherein T is $-R^{12}$ or $-alkyl-R^{12}$.
 - 29. The compound according to claim 1, wherein ${\bf R}^{12}$ is aryl or heteroaryl and is optionally substituted by 1-3 J groups.

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30. The compound according to claim 29, wherein \mathbb{R}^{12} is naphthyl, pyrazinyl, or pyridyl, any of

which is optionally substituted with a hydroxy group.

31. The compound according to claim 1, wherein ${\bf R}^{10}$ is alkyl substituted with carboxy.

- 32. The compound according to claim 1, wherein ${\bf R}^{15}$ is alkyl substituted with carboxy.
- 33. The compound according to claim 1, wherein to the ring to which R^{18} and R^{19} are attached is aromatic.
 - 34. A pharmaceutically acceptable composition comprising:
- a) a compound according to any one of claims 1
 15 33 in an amount effective to inhibit HCV NS3 protease;
 and
 - b) a pharmaceutically suitable carrier.
- 35. The use of a compound according to any one of claims 1-33 or a pharmaceutical composition according to claim 34 in the manufacture of a medicament for inhibiting serine protease activity in a patient.
 - 36. The use according to claim 35, wherein the serine protease is HCV NS3 protease.
- 37. The use of a compound according to any one of claims 1-33 or a pharmaceutical composition according to claim 34 in the manufacture of a medicament for treating or preventing a hepatitis C viral infection in a patient.

38. A process for preparing a compound of the formula (I):

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$$\stackrel{M}{\longrightarrow}$$
 $\stackrel{A^2}{\nearrow}$ $\stackrel{R^{18}}{\nearrow}$ $\stackrel{N}{\longrightarrow}$ $\stackrel{N}{$

W is:

wherein:

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m is 0 or 1;

each R^1 is hydroxy, alkoxy, or aryloxy, or each R^1 is an oxygen atom and together with the boron, to which they are each bound, form a 5-7 membered ring,

wherein the ring atoms are carbon, nitrogen, or oxygen;

each R² is independently hydrogen, alkyl, alkenyl, aryl, aralkyl, aralkenyl, cycloalkyl, cycloalkyl, cycloalkenyl, cycloalkenylalkyl, heterocyclyl, heterocyclylalkyl, heterocyclylalkenyl, heteroaryl, or heteroaralkyl, or two R² groups, which are bound to the same nitrogen atom, form together with that nitrogen atom, a 5-7 membered monocyclic heterocyclic ring system; wherein any R² carbon atom is optionally substituted with J;

J is alkyl, aryl, aralkyl, alkoxy, aryloxy, aralkoxy, cycloalkyl, cycloalkoxy, heterocyclyl, heterocyclyloxy, heterocyclylalkyl, keto, hydroxy, amino, alkylamino, alkanoylamino, aroylamino, aralkanoylamino, carboxy, carboxyalkyl, carboxamidoalkyl, halo, cyano, nitro, formyl, acyl, sulfonyl, or sulfonamido and is optionally substituted with 1-3 J¹ groups; and

J1 is alkyl, aryl, aralkyl, alkoxy, aryloxy, heterocyclyl, heterocyclyloxy, keto, hydroxy, amino, alkanoylamino, aroylamino, carboxy, carboxyalkyl, carboxamidoalkyl, halo, cyano, nitro, formyl, sulfonyl, or sulfonamido;

L is alkyl, alkenyl, or alkynyl, wherein any hydrogen is optionally substituted with halogen, and wherein any hydrogen or halogen atom bound to any terminal carbon atom is optionally substituted with sulfhydryl or hydroxy;

each M is independently alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, or

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heteroaralkyl, and is optionally substituted by 1 to 3 J groups, wherein any alkyl carbon atom may be replaced by a heteroatom;

 R^{18} is a bond, $-N(R^{11})$ -, or -C(0)-;

 R^{11} is hydrogen or C_1 - C_3 alkyl;

each R^{19} is independently H or R^{21} -aryl, or 2 adjacent R^{19} may be bound to one another to form a 5-7 membered aromatic ring; wherein any R^{19} is optionally substituted with 1 to 4 independently selected J' groups;

each R^{21} is independently C_1-C_3 -straight or branched alkyl, C_2-C_3 -straight or branched alkenyl, $O-(C_1-C_3)$ -straight or branched alkyl, or $O-(C_2-C_3)$ -straight or branched alkenyl;

n is 0 or 1;

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the ring to which R^{18} and R^{19} are attached may be saturated, partially saturated, aromatic or fully unsaturated; and 1 to 3 carbon atoms that make up the ring to which R^{18} and R^{19} are attached are optionally replaced with a heteroatom which is independently selected from O, S, S(O), S(O)₂ or N(R^{11});

 A^2 is a bond or $-N(R^{11})-R^{17}(M)-R^{22}-$, wherein

 R^{17} is -CH- or -N-; and R^{22} is -C(0)- or -S(0)₂-;

V is a bond, $-CH(R^{11})-$, -O-, -S-, or $-N(R^{11})-$;

K is a bond, -0-, -S-, -C(0)-, -S(0)-, $-S(0)_2-$, or

-S (O) NR¹¹-;

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T is $-R^{12}$, $-alkyl-R^{12}$, $-alkenyl-R^{12}$, $-alkynyl-R^{12}$, $-OR^{12}$, $-N(R^{12})_2$, $-C(O)R^{12}$, $-C(=NO-alkyl)R^{12}$, or

each R¹² is independently selected from hydrogen, aryl, heteroaryl, cycloalkyl, heterocyclyl, cycloalkylidenyl, or heterocycloalkylidenyl, and is optionally substituted with 1 to 3 J groups; or a first R¹² and a second R¹², together with the nitrogen to which they are bound, form a mono- or bicyclic ring system optionally substituted by 1 to 3 J groups;

R¹⁰ is alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, heteroaralkyl, carboxyalkyl, or carboxamidoalkyl, and is optionally substituted with 1 to 3 hydrogens J groups;

R¹⁵ is alkyl, cycloalkyl, aryl, aralkyl, heterocyclyl, heterocyclylalkyl, heteroaryl, heteroaralkyl, carboxyalkyl, or carboxamidoalkyl, and is optionally substituted with 1 to 3 J groups; and

R¹⁶ is hydrogen, alkyl, aryl, heteroaryl, cycloalkyl, or heterocyclyl; comprising the step of:

reacting a compound of formula (II):

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, wherein LG is OH or an appropriate leaving group and the other substituents are as defined above;

with a compound of formula (III):



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, wherein the NH_2 group is optionally protected and L and W are as defined above;

in the presence of a coupling reagent, provided that the compound of formula (II) or the compound of formula (III) is optionally bound to a resin.